

AMENDMENTS TO THE CLAIMS

This listing of claims replaces all prior versions, and listings, of claims in the application:

LISTING OF CLAIMS

1. (currently amended) Use of a substance for synthesising a drug for the treatment of patients who suffer from A method for inhibiting maturation of dendritic cells for the treatment of a pulmonary disease which is directly or indirectly associated to idiopathic pulmonary disease, hypersensitive pneumonia or diffused panbronchitis, wherein said substance is comprising administering to a patient a peptide or a polypeptide containing comprising the following amino acid sequence:

Arg-Lys-Gln-Met-Ala-Val-Lys-Lys-Tyr-Leu (SEQ ID NO. 4).

2. (currently amended) Use The method according to claim 1, wherein said peptide or a polypeptide further containing comprises at least one of the following amino acid sequence:

His-Ser-Asp (SEQ ID number 14); Phe-Thr-Asp (SEQ ID NO. 13).

3. (currently amended) Use The method according to claim 1, wherein said peptide or a polypeptide having following amino acid sequence:

$(A)_n$ -Arg-Lys-Gln-Met-Ala-Val-Lys-Lys-Tyr-Leu-(B)_m

wherein A, B is $(A)_n$ and $(B)_m$ independently are primary amino acid sequences comprising any sequence of natural occurring amino acid, acids; n has a value from 0 to 25 and n is the number of amino acid residues in said primary amino acid sequence

(A)_n; and m has a value from 0 to 25 and m is the number of amino acid residues in said primary amino acid sequence (B)_m. A and B are independent of each other; and n, m replacing values from 0-25; n and m are independent of each other.

4. (currently amended) Use The method according to claim 3, wherein if n > 2, said primary amino acid sequence (A)_n further comprises a primary amino acid sequence (A)_n-has following sequence if n > 2 :

(X)_o-Phe-Thr-Asp-(Y)_p;

wherein (X)_o and (Y)_p independently are primary amino acid sequences comprising X, Y is any sequence of natural occurring amino acid acids; o has a value from 0 to 11 and o is the number of amino acid residues in said primary amino acid sequence (X)_o; and p has a value from 0 to 11 and p is the number of amino acid residues in said primary amino acid sequence (Y)_p. X and Y are independently of each other; and o, p is a replacing value from 0-11, o and p are independent of each other.

5. (currently amended) Use The method according to claim 4, wherein if o > 2, said primary amino acid sequence (A)_n further comprises a primary amino acid sequence (X)_o-has following sequence if o > 2 :

(X')_q-His-Ser-Asp-(X'')_r,

wherein (X')_q and (X'')_r independently are primary amino acid sequences comprising X', X'' is any sequence of natural occurring amino acid acids; q has a value from 0 to 4 and q is the number of amino acid residues in said primary amino acid sequence (X')_q; and r has a value from 0 to 4 and r is the number of amino acid residues in said primary amino acid sequence (X'')_r. X' and X'' are independent of each other; and r, q is a replacing value from 0-4, r and q are independent of each other.

6. (currently amended) Use The method according to claim 3, wherein the sequence of said peptide or polypeptide belongs to is selected from the following group:

- (i) Arg-Lys-Gln-Met-Ala-Val-Lys-Lys-Tyr-Leu (SEQ ID NO. 4)
- (ii) Phe-Thr-Asp-X¹-X²-X³-X⁴-X⁵-Arg-Lys-Gln-Met-Ala-Val-Lys-Lys-Tyr-Leu-Asn-Ser-Ile-Leu-Asn (SEQ ID NO. 5);
- (iii) Phe-Thr-Asp-Asn-Tyr-Thr-Arg-Leu-Arg-Lys-Gln-Met-Ala-Val-Lys-Lys-Tyr-Leu-Asn-Ser-Ile-Leu-Asn (SEQ ID NO. 6);
- (iv) Phe-Thr-Asp-Ser-Tyr-Ser-Arg-Tyr-Arg-Lys-Gln-Met-Ala-Val-Lys-Lys-Tyr-Leu (SEQ ID NO. 7);
- (v) His-Ser-Asp-X¹-X²-Phe-Thr-Asp-X³-X⁴-X⁵-X⁶-X⁷-Arg-Lys-Gln-Met-Ala-Val-Lys-Lys-Tyr-Leu (SEQ ID NO. 9);
- (vi) His-Ser-Asp-Gly-Ile-Phe-Thr-Asp-Ser-Tyr-Ser-Arg-Tyr-Arg-Lys-Gln-Met-Ala-Val-Lys-Lys-Tyr-Leu (SEQ ID NO. 10);
- (vii) His-Ser-Asp-X¹-X²-Phe-Thr-Asp-Asp-X³-X⁴-X⁵-X⁶-X⁷-Arg-Lys-Gln-Met-Ala-Val-Lys-Lys-Tyr-Leu-X⁸-X⁹-X¹⁰-X¹¹-(X¹²) (SEQ ID NO. 11);
- (viii) His-Ser-Asp-Ala-Val-Phe-Thr-Asp-Asn-Tyr-Thr-Arg-Leu-Arg-Lys-Gln-Met-Ala-Val-Lys-Lys-Tyr-Leu-Asn-Ser-Ile-Leu-Asn (VIP, SEQ ID NO. 1);
- (ix) His-Ser-Asp-Gly-Ile-Phe-Thr-Asp-Ser-Tyr-Ser-Arg-Tyr-Arg-Lys-Gln-Met-Ala-Val-Lys-Lys-Tyr-Leu-Ala-Ala-Val-Leu-Gly-Lys-Arg-Tyr-Lys-Gln-Arg-Val-Lys-Asn-Lys (PACAP-38, SEQ ID NO. 2);
- (x) His-Ser-Asp-X¹-X²-Phe-Thr-Asp-X³-X⁴-X⁵-X⁶-X⁷-Arg-Lys-Gln-Met-Ala-Val-Lys-Lys-Tyr-Leu X⁸-X⁹-X¹⁰-X¹¹-X¹²-X¹³-X¹⁴-X¹⁵-X¹⁶-X¹⁷-X¹⁸-X¹⁹-X²⁰-X²¹-X²² (SEQ ID NO. 12); and
- (xi) His-Ser-Asp-Gly-Ile-Phe-Thr-Asp-Ser-Tyr-Ser-Arg-Tyr-Arg-Lys-Gln-Met-Ala-Val-Lys-Lys-Tyr-Leu-Ala-Ala-Val-Leu (PACAP-27, SEQ ID NO. 3);

and

wherein X¹ - X²² is any naturally occurring amino acid.

7. (currently amended) Use The method according to claim 1, wherein any said peptide or polypeptide is an analogue or a derivative with the same biological function.

8. (currently amended) Use The method according to claim 7, wherein any said peptide or polypeptide is in a stabilised form.

9. (currently amended) Use The method according to claim 1, wherein said disease is idiopathic pulmonary fibrosis.

10. (currently amended) Use The method according to claim 1, wherein said disease is hypersensitive pneumonia.

11. (currently amended) Use The method according to claim 1, wherein said disease is diffused panbronchiolitis.

12. (currently amended) Use The method according to claim 1, wherein the therapeutically effective peptides are administered as aerosols.

13. (currently amended) Use The method according to claim 2, wherein said disease is idiopathic pulmonary fibrosis.

14. (currently amended) Use The method according to claim 2, wherein said disease is hypersensitive pneumonia.

15. (currently amended) Use The method according to claim 2, wherein the therapeutically effective peptides are administered as aerosols.

16. (currently amended) Use The method according to claim 3, wherein any said peptide or polypeptide is an analogue or a derivative with the same biological function.

17. (currently amended) Use The method according to claim 3, wherein said disease is diffused panbronchiolitis.

18. (currently amended) Use The method according to claim 3, wherein the therapeutically effective peptides are administered as aerosols.